

**K252050 Disposable Biopsy Forceps FB-211D, FB-221D,
FB-231D, FB-241D**Mar 20, 2026
262 days to decisionK252050 · Product code: **EOB** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k252050/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nasopharyngoscope (flexible Or Rigid) (EOB)
Date received	Jul 1, 2025
Decision date	Mar 20, 2026
Days to decision	262 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Olympus Medical Systems Corporation
Location	Melville, NY, US
Contact	Seiko Yunoki
510(k) history	81 submissions · 81 cleared · 2004-2026

REGULATORY CONSULTANT

Consulting firm	Olympus Corporation of the Americas
Contact	Roshana Ahmed

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k252050/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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